## **AMENDMENTS TO THE CLAIMS**

The following listing of claims will replace all prior versions and listings of claims in the application.

## **LISTING OF CLAIMS**

What is claimed is:

- 1. (currently amended) An arteriovenous shunt comprising:
- a. an arterial graft comprising a body, a lead end and a terminal end, wherein said lead end being configured is operable for subcutaneous connection to an artery by anastomosis; and
- b. a single lumen venous outflow catheter comprising an intake end and depositing end, wherein said depositing end being configured is operable for insertion through a vein into the right atrium of the heart; and
- c. a cuff <u>operable to direct passage of blood from said arterial graft to said venous outflow catheter, said cuff</u> comprising an inlet <u>in fluid communication with</u> and outlet, wherein:
  - i. said inlet being is connected to said terminal end of said arterial subcutaneous graft; and
  - ii. said outlet being is connected to said intake end of said venous outflow catheter.
- 2. (currently amended) The arteriovenous shunt of claim 1 wherein said <u>arterial</u> subcutaneous graft is made of a biocompatible flexible material.

- 3. (original) The arteriovenous shunt of claim 2, wherein said biocompatible flexible material is polytetrafluoroethylene (PTFE) or polyurethane.
- 4. (original) The arteriovenous shunt of claim 1, wherein said arterial graft has a diameter from about 2 mm to about 8 mm and a length from about 20 cm to about 60 cm.
- 5. (original) The arteriovenous shunt of claim 4, wherein said arterial graft has a diameter of from about 6 mm to about 8 mm and a length of about 40 cm.
- 6. (original) The arteriovenous shunt of claim 1, wherein said artery is the brachial, axillary, femoral or external iliac artery.
- 7. (currently amended) The arteriovenous shunt of claim 1, wherein said cuff is polytetrafluoroethylene Teflon or polyethylene terephthalate Dacron.
- 8. (original) The arteriovenous shunt of claim 1, wherein said venous outflow catheter has a diameter from about 1 mm to about 7 mm and a length of from about 20 cm to about 80 cm.
- 9. (original) The arteriovenous shunt of claim 1, wherein said venous outflow catheter has a diameter from about 5 mm to about 7 mm and a length of from about 40 cm to about 60 cm.

- 10. (original) The arteriovenous shunt of claim 1, wherein said venous outflow catheter is made of polyurethane or silicone.
- 11. (original) The arteriovenous shunt of claim 1, wherein said vein is the cephalic, axillary, jugular, femoral or external iliac vein.
- 12. (currently amended) The arteriovenous shunt of claim 1, wherein said venous outflow catheter has a diameter of about 1 mm smaller than said arterial the subcutaneous graft.
- 13. (currently amended) A system for performing hemodialysis on a patient comprising:
  - a. an arteriovenous shunt comprising:
    - i. an arterial graft comprising a body, a lead end and a terminal end,
       wherein said lead end being configured is operable for subcutaneous connection to an artery by anastomosis; and
    - ii. a single lumen venous outflow catheter comprising an intake end and depositing end, wherein said depositing end being configured is operable for insertion through a vein into the right atrium of the heart; and
    - iii. ii. a cuff operable to direct passage of blood from said arterial graft to said venous outflow catheter, said cuff comprising an inlet in fluid communication with and an outlet, wherein:

- said inlet <u>being</u> is connected to said terminal end of said subcutaneous graft; and
- said outlet <u>being</u> is connected to said intake end of said venous outflow catheter;

and

- b. a hemodialysis apparatus.
- 14. (currently amended) The system according to claim 13, wherein said venous outflow catheter has a diameter of about 1 mm smaller than said <u>arterial</u> subcutaneous graft.
- 15. (original) The system according to claim 13, wherein said artery is the brachial, axillary, femoral or external iliac artery.
- 16. (original) The system according to claim 13, wherein said vein is the cephalic, axillary, jugular, femoral or external iliac vein.
- 17. (currently amended) A method of performing hemodialysis on a patient comprising:
- a. <u>surgically</u> inserting an arteriovenous shunt into a patient, wherein said arteriovenous shunt comprises:
  - i. an arterial graft comprising a body, a lead end and a terminal end,

    wherein said lead end being configured is operable for subcutaneous

    connection to an artery by anastomosis; and

- ii. a single lumen venous outflow catheter comprising an intake end and depositing end, wherein said depositing end being configured is operable for insertion through a vein into the right atrium of the heart; and
- iii. a cuff operable to direct passage of blood from said arterial graft to said venous outflow catheter, said cuff comprising an inlet in fluid communication with and an outlet, wherein:
  - said inlet <u>being</u> is connected to said terminal end of said <u>arterial</u>
     subcutaneous graft; and
  - said outlet <u>being</u> is connected to said intake end of said venous outflow catheter;
- b. connecting said arterial graft to a hemodialysis apparatus;
- c. collecting blood from the patient through said arterial subcutaneous graft;
- d. passing said blood through the hemodialysis apparatus;
- e. collecting purified blood from hemodialysis apparatus; and
- f. transmitting said purified blood through said cuff into said venous outflow catheter.
- 18. (currently amended) The method according to claim 16 wherein said venous outflow catheter has a diameter of about 1 mm smaller than said <u>arterial</u> subcutaneous graft.
- 19. (original) The method according to claim 16, wherein said artery is the brachial, axillary, or femoral, external iliac artery.

	20.	(original) The method according to claim 16, wherein said vein is the axillary
jugula	ar, femo	oral or external iliac vein.